

Wolfgang Hogler

List of Publications by Year in descending order

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Version: 2024-02-01

137
papers

7,156
citations

66234

42
h-index

64668

79
g-index

143
all docs

143
docs citations

143
times ranked

6931
citing authors

#	ARTICLE	IF	CITATIONS
1	Breast satisfaction in adult women with Turner syndrome—An international survey employing the BREAST-Q questionnaire. <i>Clinical Endocrinology</i> , 2023, 98, 82-90.	1.2	3
2	Sustained Efficacy and Safety of Burosumab, a Monoclonal Antibody to FGF23, in Children With X-Linked Hypophosphatemia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, 813-824.	1.8	36
3	Osteoporosis in children and adolescents: when to suspect and how to diagnose it. <i>European Journal of Pediatrics</i> , 2022, 181, 2549-2561.	1.3	24
4	Effect of Burosumab Compared With Conventional Therapy on Younger vs Older Children With X-linked Hypophosphatemia. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2022, 107, e3241-e3253.	1.8	36
5	Spontaneous reshaping of vertebral fractures in an adolescent with osteogenesis imperfecta. <i>Bone Reports</i> , 2022, 16, 101595.	0.2	3
6	The landscape of retesting in childhood-onset idiopathic growth hormone deficiency and its reversibility: a systematic review and meta-analysis. <i>European Journal of Endocrinology</i> , 2022, 187, 265-278.	1.9	6
7	SMAD3 mutation in LDS3 causes bone fragility by impairing the TGF- β 2 pathway and enhancing osteoclastogenesis. <i>Bone Reports</i> , 2022, 17, 101603.	0.2	3
8	Failure of national antenatal vitamin D supplementation programme puts dark skinned infants at highest risk: A newborn bloodspot screening study. <i>Clinical Nutrition</i> , 2021, 40, 3542-3551.	2.3	11
9	Mechanisms of Bone Fragility: From Osteogenesis Imperfecta to Secondary Osteoporosis. <i>International Journal of Molecular Sciences</i> , 2021, 22, 625.	1.8	44
10	Patient-Reported Outcomes from a Randomized, Active-Controlled, Open-Label, Phase 3 Trial of Burosumab Versus Conventional Therapy in Children with X-Linked Hypophosphatemia. <i>Calcified Tissue International</i> , 2021, 108, 622-633.	1.5	26
11	The effect of vitamin D supplementation and nutritional intake on skeletal maturity and bone health in socio-economically deprived children. <i>European Journal of Nutrition</i> , 2021, 60, 3343-3353.	1.8	3
12	Recombinant human parathyroid hormone (1-84) replacement therapy in a child with hypoparathyroidism. <i>Bone</i> , 2021, 144, 115834.	1.4	1
13	Activating mutations in BRAF disrupt the hypothalamo-pituitary axis leading to hypopituitarism in mice and humans. <i>Nature Communications</i> , 2021, 12, 2028.	5.8	12
14	Growth and Body Composition in PKU Children—A Three-Year Prospective Study Comparing the Effects of L-Amino Acid to Glycomacropeptide Protein Substitutes. <i>Nutrients</i> , 2021, 13, 1323.	1.7	12
15	A Three-Year Longitudinal Study Comparing Bone Mass, Density, and Geometry Measured by DXA, pQCT, and Bone Turnover Markers in Children with PKU Taking L-Amino Acid or Glycomacropeptide Protein Substitutes. <i>Nutrients</i> , 2021, 13, 2075.	1.7	7
16	A Roadmap to Gene Discoveries and Novel Therapies in Monogenic Low and High Bone Mass Disorders. <i>Frontiers in Endocrinology</i> , 2021, 12, 709711.	1.5	13
17	Case Report: Severe Neonatal Course in Paternally Derived Familial Hypocalciuric Hypercalcemia. <i>Frontiers in Endocrinology</i> , 2021, 12, 700612.	1.5	3
18	A novel cryptic splice site mutation in COL1A2 as a cause of osteogenesis imperfecta. <i>Bone Reports</i> , 2021, 15, 101110.	0.2	3

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19	Bone density and body composition in small for gestational age children with adequate catch up growth: A preliminary retrospective case control study. <i>Bone</i> , 2021, 153, 116114.	1.4	3
20	Bone Phenotyping Approaches in Human, Mice and Zebrafish – Expert Overview of the EU Cost Action GEMSTONE (Genomics of MusculoSkeletal traits Translational Network). <i>Frontiers in Endocrinology</i> , 2021, 12, 720728.	1.5	12
21	Micronutrient deficiencies and health-related quality of life: the case of children with vitamin D deficiency. <i>Public Health Nutrition</i> , 2020, 23, 1165-1172.	1.1	7
22	Diagnosis, treatment-monitoring and follow-up of children and adolescents with X-linked hypophosphatemia (XLH). <i>Metabolism: Clinical and Experimental</i> , 2020, 103, 153892.	1.5	46
23	Guidance to Bone Morbidity in Children and Adolescents Undergoing Allogeneic Hematopoietic Stem Cell Transplantation. <i>Biology of Blood and Marrow Transplantation</i> , 2020, 26, e27-e37.	2.0	6
24	Oral Iron for Prevention and Treatment of Rickets and Osteomalacia in Autosomal Dominant Hypophosphatemia. <i>Journal of Bone and Mineral Research</i> , 2020, 35, 226-230.	3.1	6
25	A Contemporary View of the Definition and Diagnosis of Osteoporosis in Children and Adolescents. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2020, 105, e2088-e2097.	1.8	64
26	Burden of Illness in Adults With Hypophosphatasia: Data From the Global Hypophosphatasia Patient Registry. <i>Journal of Bone and Mineral Research</i> , 2020, 35, 2171-2178.	3.1	38
27	Vitamin D supplementation in pregnancy, lactation and infancy: why is it fundamental?. <i>British Journal of Midwifery</i> , 2020, 28, 315-322.	0.1	0
28	Prevention of vitamin D deficiency improves population health, social inequalities and health care budgets. <i>European Journal of Public Health</i> , 2020, 30, 393-394.	0.1	2
29	New developments in the management of achondroplasia. <i>Wiener Medizinische Wochenschrift</i> , 2020, 170, 104-111.	0.5	18
30	New Developments in the Treatment of X-Linked Hypophosphataemia: Implications for Clinical Management. <i>Paediatric Drugs</i> , 2020, 22, 113-121.	1.3	12
31	Nutritional rickets & osteomalacia: A practical approach to management. <i>Indian Journal of Medical Research</i> , 2020, 152, 356.	0.4	27
32	Biallelic KIF24 Variants Are Responsible for a Spectrum of Skeletal Disorders Ranging From Lethal Skeletal Ciliopathy to Severe Acromesomelic Dysplasia. <i>Journal of Bone and Mineral Research</i> , 2020, 37, 1642-1652.	3.1	5
33	The Utility of DXA Assessment at the Forearm, Proximal Femur, and Lateral Distal Femur, and Vertebral Fracture Assessment in the Pediatric Population: 2019 ISCD Official Position. <i>Journal of Clinical Densitometry</i> , 2019, 22, 567-589.	0.5	83
34	Efficacy and Safety of Asfotase Alfa in Infants and Young Children With Hypophosphatasia: A Phase 2 Open-Label Study. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2019, 104, 2735-2747.	1.8	46
35	Tissue non-specific alkaline phosphatase activity and mineralization capacity of bi-allelic mutations from severe perinatal and asymptomatic hypophosphatasia phenotypes: Results from an in vitro mutagenesis model. <i>Bone</i> , 2019, 127, 9-16.	1.4	8
36	Older age at initiation of antiretroviral therapy predicts low bone mineral density in children with perinatally-infected HIV in Zimbabwe. <i>Bone</i> , 2019, 125, 96-102.	1.4	19

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37	Burosumab versus conventional therapy in children with X-linked hypophosphataemia: a randomised, active-controlled, open-label, phase 3 trial. <i>Lancet, The</i> , 2019, 393, 2416-2427.	6.3	229
38	Bone density in children: what are we measuring?. <i>Archives of Disease in Childhood</i> , 2019, 104, 1108-1111.	1.0	8
39	Diagnostic delay is common among patients with hypophosphatasia: initial findings from a longitudinal, prospective, global registry. <i>BMC Musculoskeletal Disorders</i> , 2019, 20, 80.	0.8	69
40	Rickets and Osteomalacia. , 2019, , 339-354.		2
41	Spot the silent sufferers: A call for clinical diagnostic criteria for solar and nutritional osteomalacia. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2019, 188, 141-146.	1.2	30
42	Causes, patterns and severity of androgen excess in 487 consecutively recruited pre- and post-pubertal children. <i>European Journal of Endocrinology</i> , 2019, 180, 213-221.	1.9	22
43	Comparing modalities of conducting the six-minute walk test in healthy children and adolescents. <i>Minerva Pediatrica</i> , 2019, 71, 229-234.	2.6	5
44	OR13-4 Safety Profile of Asfotase Alfa Treatment of Patients with Hypophosphatasia: A Pooled Analysis. <i>Journal of the Endocrine Society</i> , 2019, 3, .	0.1	0
45	Prevention of rickets and osteomalacia in the UK: political action overdue. <i>Archives of Disease in Childhood</i> , 2018, 103, 901-906.	1.0	46
46	Osteonecrosis of the Jaw and Rebound Hypercalcemia in Young People Treated With Denosumab for Giant Cell Tumor of Bone. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2018, 103, 596-603.	1.8	56
47	Efficacy and treatment costs of zoledronate versus pamidronate in paediatric osteoporosis. <i>Archives of Disease in Childhood</i> , 2018, 103, 92-94.	1.0	16
48	Continuous Subcutaneous Recombinant Parathyroid Hormone (1â€“34) Infusion in the Management of Childhood Hypoparathyroidism Associated with Malabsorption. <i>Hormone Research in Paediatrics</i> , 2018, 89, 271-277.	0.8	24
49	Burosumab Therapy in Children with X-Linked Hypophosphatemia. <i>New England Journal of Medicine</i> , 2018, 378, 1987-1998.	13.9	339
50	Bone Mineral Density Corrected for Size in Childhood Leukaemia Survivors Treated with Haematopoietic Stem Cell Transplantation and Total Body Irradiation. <i>Hormone Research in Paediatrics</i> , 2018, 89, 246-254.	0.8	3
51	Cumulative radiation exposure from medical imaging and associated lifetime cancer risk in children with osteogenesis imperfecta. <i>Bone</i> , 2018, 114, 252-256.	1.4	10
52	Growth, bone health & ambulatory status of boys with DMD treated with daily vs. intermittent oral glucocorticoid regimen. <i>Bone</i> , 2018, 116, 181-186.	1.4	28
53	Cardiac, bone and growth plate manifestations in hypocalcemic infants: revealing the hidden body of the vitamin D deficiency iceberg. <i>BMC Pediatrics</i> , 2018, 18, 183.	0.7	43
54	Bone Density in the Obese Child: Clinical Considerations and Diagnostic Challenges. <i>Calcified Tissue International</i> , 2017, 100, 514-527.	1.5	25

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55	Vertebral fractures assessment in children: Evaluation of DXA imaging versus conventional spine radiography. <i>Bone</i> , 2017, 97, 168-174.	1.4	44
56	The Effect of Whole Body Vibration Training on Bone and Muscle Function in Children With Osteogenesis Imperfecta. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 2734-2743.	1.8	28
57	Phenotypic Spectrum in Osteogenesis Imperfecta Due to Mutations in TMEM38B: Unraveling a Complex Cellular Defect. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 2019-2028.	1.8	27
58	Hormone supplementation for pubertal induction in girls. <i>Archives of Disease in Childhood</i> , 2017, 102, 975-980.	1.0	47
59	Nutritional Rickets and Osteomalacia in the Twenty-first Century: Revised Concepts, Public Health, and Prevention Strategies. <i>Current Osteoporosis Reports</i> , 2017, 15, 293-302.	1.5	97
60	Variations in infant and childhood vitamin D supplementation programmes across Europe and factors influencing adherence. <i>Endocrine Connections</i> , 2017, 6, 667-675.	0.8	57
61	Preventing vitamin D deficiency (VDD): a systematic review of economic evaluations. <i>European Journal of Public Health</i> , 2017, 27, 292-301.	0.1	24
62	Amalgamated Reference Data for Size-Adjusted Bone Densitometry Measurements in 3598 Children and Young Adults—the ALPHABET Study. <i>Journal of Bone and Mineral Research</i> , 2017, 32, 172-180.	3.1	98
63	Bone Structural Characteristics and Response to Bisphosphonate Treatment in Children With Hajdu-Cheney Syndrome. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 4163-4172.	1.8	22
64	Response to Letter to the Editor: “The Effect of Whole Body Vibration Training on Bone and Muscle Function in Children With Osteogenesis Imperfecta”. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2017, 102, 4262-4263.	1.8	2
65	Absence of the ER Cation Channel TMEM38B/TRIC-B Disrupts Intracellular Calcium Homeostasis and Dysregulates Collagen Synthesis in Recessive Osteogenesis Imperfecta. <i>PLoS Genetics</i> , 2016, 12, e1006156.	1.5	49
66	Recurrent Proximal Femur Fractures in a Teenager With Osteogenesis Imperfecta on Continuous Bisphosphonate Therapy: Are We Overtreating?. <i>Journal of Bone and Mineral Research</i> , 2016, 31, 1449-1454.	3.1	33
67	Rickets and osteomalacia: a call for action to protect immigrants and ethnic risk groups. <i>The Lancet Global Health</i> , 2016, 4, e229-e230.	2.9	18
68	Natural history of retinopathy in children and young people with type 1 diabetes. <i>Eye</i> , 2016, 30, 987-991.	1.1	8
69	Phenotypic variability in patients with osteogenesis imperfecta caused by <i>BMP1</i> mutations. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 3150-3156.	0.7	32
70	Nutritional rickets in immigrant and refugee children. <i>Public Health Reviews</i> , 2016, 37, 3.	1.3	55
71	Global Consensus Recommendations on Prevention and Management of Nutritional Rickets. <i>Hormone Research in Paediatrics</i> , 2016, 85, 83-106.	0.8	158
72	A Diagnostic Algorithm for Children with Low Alkaline Phosphatase Activities: Lessons Learned from Laboratory Screening for Hypophosphatasia. <i>Journal of Pediatrics</i> , 2016, 172, 181-186.e1.	0.9	33

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73	Global Consensus Recommendations on Prevention and Management of Nutritional Rickets. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 394-415.	1.8	774
74	Consensus Recommendations for Prevention of Nutritional Rickets: Food Fortification and Micronutrient Supplements for Global Health. <i>AIMS Public Health</i> , 2016, 3, 40-48.	1.1	6
75	Complications of vitamin D deficiency from the foetus to the infant: One cause, one prevention, but who's responsibility?. <i>Best Practice and Research in Clinical Endocrinology and Metabolism</i> , 2015, 29, 385-398.	2.2	49
76	ENDOCRINOLOGY AND ADOLESCENCE: Osteoporosis in children: diagnosis and management. <i>European Journal of Endocrinology</i> , 2015, 173, R185-R197.	1.9	64
77	Sex-, age-, and height-specific reference curves for the 6-min walk test in healthy children and adolescents. <i>European Journal of Pediatrics</i> , 2015, 174, 837-840.	1.3	29
78	Osteoporosis in Children with Chronic Disease. <i>Endocrine Development</i> , 2015, 28, 176-195.	1.3	23
79	Recombinant Human Growth Hormone Therapy in Children with Chromosome 15q26 Deletion. <i>Hormone Research in Paediatrics</i> , 2015, 83, 424-430.	0.8	6
80	Iron Supplementation Associated With Loss of Phenotype in Autosomal Dominant Hypophosphatemic Rickets. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2015, 100, 3388-3392.	1.8	40
81	The modulation of corticosteroid metabolism by hydrocortisone therapy in patients with hypopituitarism increases tissue glucocorticoid exposure. <i>European Journal of Endocrinology</i> , 2015, 173, 583-593.	1.9	13
82	The relationship between bone mass and body composition in children with hypothalamic and simple obesity. <i>Clinical Endocrinology</i> , 2014, 80, 85-91.	1.2	7
83	IGFALS Gene Dosage Effects on Serum IGF-I and Glucose Metabolism, Body Composition, Bone Growth in Length and Width, and the Pharmacokinetics of Recombinant Human IGF-I Administration. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2014, 99, E703-E712.	1.8	25
84	Longitudinal changes in glucocorticoid metabolism are associated with later development of adverse metabolic phenotype. <i>European Journal of Endocrinology</i> , 2014, 171, 433-442.	1.9	24
85	Bone Health in Children and Adolescents With Chronic Diseases That May Affect the Skeleton: The 2013 ISCD Pediatric Official Positions. <i>Journal of Clinical Densitometry</i> , 2014, 17, 281-294.	0.5	119
86	Thyroid dysmorphogenesis is mainly caused by <i>TPO</i> mutations in consanguineous community. <i>Clinical Endocrinology</i> , 2013, 79, 275-281.	1.2	47
87	Diagnostic evaluation of bone densitometric size adjustment techniques in children with and without low trauma fractures. <i>Osteoporosis International</i> , 2013, 24, 2015-2024.	1.3	58
88	Relationship between serum 25-hydroxyvitamin D and parathyroid hormone in the search for a biochemical definition of vitamin D deficiency in children. <i>Pediatric Research</i> , 2013, 74, 552-556.	1.1	95
89	Lower CD28+ T cell proportions were associated with CMV-seropositivity in patients with Hashimoto's thyroiditis. <i>BMC Endocrine Disorders</i> , 2013, 13, 34.	0.9	8
90	A Diagnosis Not to Be Missed: Nonclassic Steroid 11 β -Hydroxylase Deficiency Presenting With Premature Adrenarche and Hirsutism. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2013, 98, E1620-E1625.	1.8	63

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91	Five novel mutations in the SCNN1A gene causing autosomal recessive pseudohypoaldosteronism type 1. <i>European Journal of Endocrinology</i> , 2013, 168, 707-715.	1.9	25
92	Clinical characteristics and management of cranial diabetes insipidus in infants. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2013, 26, 1041-6.	0.4	16
93	An association of craniopharyngioma in Turner syndrome. <i>Pediatric Blood and Cancer</i> , 2013, 60, E7-E9.	0.8	6
94	TSHR is the main causative locus in autosomal recessively inherited thyroid dysgenesis. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2012, 25, 419-26.	0.4	27
95	Endocrine and Bone Metabolic Complications in Chronic Liver Disease and After Liver Transplantation in Children. <i>Journal of Pediatric Gastroenterology and Nutrition</i> , 2012, 54, 313-321.	0.9	59
96	Whole-exome sequencing identifies mutations in the nucleoside transporter gene SLC29A3 in dysosteosclerosis, a form of osteopetrosis. <i>Human Molecular Genetics</i> , 2012, 21, 4904-4909.	1.4	81
97	Biochemical Markers of Bone Metabolism. , 2012, , 361-381.		126
98	Salt-losing crisis in infantsâ€”not always of adrenal origin. <i>European Journal of Pediatrics</i> , 2012, 171, 317-321.	1.3	14
99	Six-Minute Walk Distance in Overweight Children and Adolescents: Effects of a Weight-Reducing Program. <i>Journal of Pediatrics</i> , 2011, 158, 447-451.	0.9	30
100	Galactokinase Deficiency in a Patient with Congenital Hyperinsulinism. <i>JIMD Reports</i> , 2011, 5, 7-11.	0.7	3
101	Fractures and Fanconi Syndrome due to Prolonged Sodium Valproate Use. <i>Neuropediatrics</i> , 2011, 42, 119-121.	0.3	23
102	Chronic antiepileptic monotherapy, bone metabolism, and body composition in nonâ€”institutionalized children. <i>Developmental Medicine and Child Neurology</i> , 2010, 52, 283-288.	1.1	49
103	Childhood growth hormone deficiency, bone density, structures and fractures: scrutinizing the evidence. <i>Clinical Endocrinology</i> , 2010, 72, 281-289.	1.2	78
104	Novel <i>TSHR</i> mutations in consanguineous families with congenital nongoitrous hypothyroidism. <i>Clinical Endocrinology</i> , 2010, 73, 671-677.	1.2	28
105	Z-Score Comparability of Bone Mineral Density Reference Databases for Children. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 4652-4659.	1.8	35
106	Impact of Heterozygosity for Acid-Labile Subunit (IGFALS) Gene Mutations on Stature: Results from the International Acid-Labile Subunit Consortium. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2010, 95, 4184-4191.	1.8	52
107	Oxcarbazepine accelerates cortisol elimination via cytochrome P450 3A4 induction. <i>Archives of Disease in Childhood</i> , 2010, 95, 1065-1065.	1.0	22
108	Growth and bone health in chronic liver disease and following liver transplantation in children. <i>Pediatric Endocrinology Reviews</i> , 2010, 7, 266-74.	1.2	21

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109	Three Novel & i>>IGFALS</i> Gene Mutations Resulting in Total ALS and Severe Circulating IGF-I/IGFBP-3 Deficiency in Children of Different Ethnic Origins. <i>Hormone Research in Paediatrics</i> , 2009, 71, 100-110.	0.8	29
110	Thoughts on paediatric diabetes care in the UK. <i>British Journal of Diabetes and Vascular Disease</i> , 2009, 9, 259-267.	0.6	4
111	Bone mineral density in young adult survivors of acute lymphoblastic leukemia. <i>Cancer</i> , 2009, 115, 4885-4885.	2.0	6
112	PPIB Mutations Cause Severe Osteogenesis Imperfecta. <i>American Journal of Human Genetics</i> , 2009, 85, 521-527.	2.6	257
113	Pediatric reference intervals for thyroid hormone levels from birth to adulthood: a retrospective study. <i>BMC Endocrine Disorders</i> , 2008, 8, 15.	0.9	137
114	Sex-specific developmental changes in muscle size and bone geometry at the femoral shaft. <i>Bone</i> , 2008, 42, 982-989.	1.4	38
115	Dual Energy X-ray Absorptiometry Interpretation and Reporting in Children and Adolescents: The 2007 ISCD Pediatric Official Positions. <i>Journal of Clinical Densitometry</i> , 2008, 11, 43-58.	0.5	480
116	Dual-Energy X-ray Aborptiometry Assessment in Children and Adolescents with Diseases that May Affect the Skeleton: The 2007 ISCD Pediatric Official Positions. <i>Journal of Clinical Densitometry</i> , 2008, 11, 29-42.	0.5	104
117	Sex- and Age-Specific Reference Curves for Serum Markers of Bone Turnover in Healthy Children from 2 Months to 18 Years. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2007, 92, 443-449.	1.8	231
118	Pyridoxine-responsive seizures as the first symptom of infantile hypophosphatasia caused by two novel missense mutations (c.677T>C, p.M226T; c.1112C>T, p.T371I) of the tissue-nonspecific alkaline phosphatase gene. <i>Bone</i> , 2007, 40, 1655-1661.	1.4	141
119	Acute phase response and mineral status following low dose intravenous zoledronic acid in children. <i>Bone</i> , 2007, 41, 366-370.	1.4	75
120	Incidence of skeletal complications during treatment of childhood acute lymphoblastic leukemia: Comparison of fracture risk with the General Practice Research Database. <i>Pediatric Blood and Cancer</i> , 2007, 48, 21-27.	0.8	91
121	Brain-type Natriuretic Peptide Release and Seizure Activity during Vagal Nerve Stimulation. <i>Epilepsia</i> , 2007, 48, 397-399.	2.6	15
122	B-cell lymphoma in a girl with ataxia teleangiectasia (A-T) treated with rituximab monotherapy. <i>Pediatric Blood and Cancer</i> , 2006, 46, 528-529.	0.8	16
123	The re-emerging burden of rickets: a decade of experience from Sydney. <i>Archives of Disease in Childhood</i> , 2005, 91, 564-568.	1.0	169
124	Effect of growth hormone therapy and puberty on bone and body composition in children with idiopathic short stature and growth hormone deficiency. <i>Bone</i> , 2005, 37, 642-650.	1.4	66
125	Relation between hormones and body composition, including bone, in prepubertal children. <i>American Journal of Clinical Nutrition</i> , 2004, 80, 966-972.	2.2	173
126	Importance of Estrogen on Bone Health in Turner Syndrome: A Cross-Sectional and Longitudinal Study Using Dual-Energy X-Ray Absorptiometry. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004, 89, 193-199.	1.8	97

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127	The Skeletal Phenotype of Men with Previous Constitutional Delay of Puberty. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2004, 89, 4306-4311.	1.8	53
128	Severe transient hyperinsulinaemic hypoglycaemia: two neonates without predisposing factors and a review of the literature. <i>European Journal of Pediatrics</i> , 2004, 163, 38-41.	1.3	28
129	Short-term safety assessment in the use of intravenous zoledronic acid in children. <i>Journal of Pediatrics</i> , 2004, 145, 701-704.	0.9	67
130	Importance of lean mass in the interpretation of total body densitometry in children and adolescents. <i>Journal of Pediatrics</i> , 2003, 143, 81-88.	0.9	162
131	A comparison of bone geometry and cortical density at the mid-femur between prepuberty and young adulthood using magnetic resonance imaging. <i>Bone</i> , 2003, 33, 771-778.	1.4	71
132	Perinatal Bone Turnover in Term Human Neonates and the Influence of Maternal Smoking. <i>Pediatric Research</i> , 2003, 53, 817-822.	1.1	17
133	Rhinocerebral Mucormycosis in a Boy With Recurrent Acute Lymphoblastic Leukemia: Long-Term Survival With Systemic Antifungal Treatment. <i>Journal of Pediatric Hematology/Oncology</i> , 2002, 24, 492-494.	0.3	18
134	Abrupt onset of severe passenger lymphocyte syndrome after HLA-mismatched unrelated bone marrow transplantation in a seven-year-old boy. <i>Medical and Pediatric Oncology</i> , 2002, 38, 143-144.	1.0	5
135	Prolonged iron depletion after allogeneic 2-unit RBC apheresis. <i>Transfusion</i> , 2001, 41, 602-605.	0.8	23
136	Characteristics of Vitamin D supplementation programs for the prevention of rickets in infants and young children in Europe: Factors influencing compliance. <i>Endocrine Abstracts</i> , 0, , .	0.0	1
137	Effect of KRN23, a fully human anti-FGF23 monoclonal antibody, on rickets in children with X-linked hypophosphatemia (XLH): 40-week interim results from a randomized, open-label Phase 2 study. <i>Bone Abstracts</i> , 0, , .	0.0	0